

**Amendments to the Claims:**

1. (Currently Amended) A method of labeling a ~~molecule~~ protein exposed on a luminal surface of a cell lining of a perfusable space *in situ* or *in vivo* comprising the following steps:
  - (a) providing a cell membrane impermeable reagent comprising three domains
    - (i) a first domain comprising a chemical moiety capable of covalently and non-specifically binding to ~~a molecule~~ the protein exposed on the luminal surface of a cell lining of a perfusable space *in situ* or *in vivo*,
    - (ii) a second domain comprising a labeling domain, and
    - (iii) a third domain situated between the first and second domains linking the first domain to the second domain by a cleavable chemical moiety, wherein the cleavable chemical moiety ~~is not cleavable under *in vivo* conditions but is cleavable under a condition that does not denature the lumen-exposed molecule~~ comprises a disulfide group, a periodate-cleavable glycol, a dithionite-cleavable diazobond, a hydroxylamine-cleavable ester, or a base-labile sulfone;
  - (b) administering the membrane impermeable reagent into the perfusable space in an intact organ or an intact animal to react the cell membrane impermeable reagent with ~~the molecule~~ proteins expressed exposed on the luminal surface of the cell lining of the perfusable space to label the lumen-exposed ~~molecule~~ protein; and
  - (c) cleaving the cleavable chemical moiety of the reagent that reacted with the lumen exposed ~~molecule~~ protein under ~~the~~ a condition that does not denature the lumen-exposed ~~molecule~~ protein.
2. (Currently Amended) The method of claim 1, wherein the reagent-reacted lumen-exposed ~~molecule~~ protein is an organ-specific or a tissue-specific ~~molecule~~ protein.
3. (Original) The method of claim 1, wherein the perfusable space is a lumen of a vascular vessel and the cell lining the space is an endothelial cell.

4. (Original) The method of claim 3, wherein the vascular vessel is an artery, an arteriole, a vein, or a capillary.
5. (Original) The method of claim 1, wherein the perfusable space is a lumen of a cerebral spinal fluid (CSF) space.
6. (Original) The method of claim 1, wherein the perfusable space is a lumen of a lymphatic vessel and the cell lining the space is an endothelial cell.
7. (Original) The method of claim 1, wherein the perfusable space is a lumen of an endocrine or exocrine duct or pore.
8. (Original) The method of claim 1, wherein the cell lining the perfusable space is an epithelial cell.
9. (Original) The method of claim 1, wherein the organ is, or the tissue is derived from, a heart, a lung, a brain, a liver, a kidney, an endocrine gland, skin, a reproductive organ, a digestive tract organ, or an eye.
10. (Previously Presented) The method of claim 1, wherein the labeling domain of the reagent is selected from the group consisting of an enzyme, biotin, a colorimetric moiety, a fluorescent moiety, a luminescent moiety, a bioluminescent moiety, a radionucleotide and a paramagnetic element.
- 11-12. (Cancelled)
13. (Original) The method of claim 1, wherein the cleavable chemical moiety comprises a disulfide group.
- 14-15. (Cancelled)

16. (Original) The method of claim 1, wherein administering the cell membrane impermeable reagent into the perfusable space of the intact organ or tissue or the intact animal comprises administration of a buffered, aqueous solution comprising the cell membrane impermeable reagent.

17-18. (Cancelled)

19. (Currently Amended) A method of isolating a ~~molecule~~ protein that is exposed on a luminal surface of a perfusable space comprising the following steps:

- (a) providing a cell membrane impermeable reagent comprising three domains
  - (i) a first domain comprising a chemical moiety capable of covalently and non-specifically binding to a ~~molecule~~ protein expressed on the luminal surface of a cell lining a perfusable space *in situ* or *in vivo*,
  - (ii) a second domain comprising a binding domain;
  - (iii) a third domain situated between the first and second domains linking the first domain to the second domain by a cleavable chemical moiety, wherein the cleavable chemical moiety ~~is not cleavable under *in vivo* conditions but is cleavable under a condition that does not denature the lumen-exposed molecule~~ comprises a disulfide group, a periodate-cleavable glycol, a dithionite-cleavable diazobond, a hydroxylamine-cleavable ester, or a base-labile sulfone;
- (b) administering the cell membrane impermeable reagent into the perfusable space in an intact organ or an intact animal to react the cell membrane impermeable reagent with a ~~molecule-protein~~ expressed exposed on the luminal surface of the cell lining of the perfusable space; and
- (c) cleaving the cleavable chemical moiety of the reagent that reacted with the lumen exposed protein under a condition that does not denature the lumen-exposed protein;  
and
- (d) isolating the reagent-reacted lumen-exposed ~~molecule-protein~~ by contacting the reagent-reacted lumen-exposed protein with a ligand that reacted with the reagent under the a condition that does not denature the lumen-exposed molecule.

20. (Currently Amended) The method of claim 19, wherein the lumen-exposed ~~molecule~~ protein is an organ-specific or a tissue-specific ~~molecule~~ protein.
21. (Currently Amended) The method of claim 20, further comprising the step of comparing the reagent-reacted ~~molecules~~ proteins from different organs or tissues to identify the organ-specific or tissue-specific ~~molecule~~ protein, wherein the organ-specific or tissue-specific ~~molecule~~ protein is exposed on the luminal surface of the perfusable space of only one of the compared organs or tissues.
22. (Original) The method of claim 19, wherein the perfusable space is a lumen of a vascular vessel and the cell lining the space is an endothelial cell.
23. (Original) The method of claim 22, wherein the vascular vessel is an artery, an arteriole, a vein, or a capillary.
24. (Original) The method of claim 19, wherein the perfusable space is a lumen of a cerebral spinal fluid (CSF) space.
25. (Original) The method of claim 19, wherein the perfusable space is a lumen of a lymphatic vessel and the cell lining the space is an endothelial cell.
26. (Original) The method of claim 19, wherein the perfusable space is a lumen of an endocrine or exocrine duct or pore.
27. (Currently Amended) The method of claim 19, wherein the cell lining of the perfusable space is an epithelial cell.
28. (Original) The method of claim 19, wherein the organ is, or the tissue is derived from, a heart, a lung, a brain, a liver, a kidney, an endocrine gland, skin, a reproductive organ, a digestive tract organ, or an eye.

29. (Currently Amended) The method of claim 19, wherein the ligand ~~binding domain of the reagent~~ comprises of biotin.

30. (Currently Amended) The method of claim 19, wherein the ligand ~~binding domain of the reagent~~ comprises a polypeptide, a nucleic acid, or a peptide nucleic acid.

31. (Withdrawn) The method of claim 30, wherein the polypeptide comprises a polyhistidine, a protein A domain, or a FLAG extension.

32. (Original) The method of claim 19, wherein the cleavable chemical moiety comprises a disulfide group.

33-35. (Cancelled)

36. (Original) The method of claim 19, wherein administering the cell membrane impermeable reagent into the perfusable space of the intact organ or tissue or the intact animal comprises administration of a buffered, aqueous solution comprising the cell membrane impermeable reagent.

37-38. (Cancelled)

39. (Original) The method of claim 19, wherein two separate cell membrane impermeable reagents are co-administered.

40. (Currently Amended) The method of claim 19, wherein the reagent-reacted ~~molecule~~ protein is isolated by

- (a) contacting a cell or a membrane isolate or a cell or a tissue homogenate or an extract derived from the reagent-reacted organ or animal with a the ligand having affinity for the binding domain of the cell membrane impermeable reagent; and
- (b) removing a non-bound ~~molecule~~ protein from the ligand-bound ~~molecules~~ proteins.

41. (Original) The method of claim 40, wherein the ligand is immobilized.
42. (Original) The method of claim 41, wherein the ligand is immobilized on a bead.
43. (Currently Amended) The method of claim 40, wherein the ~~binding-domain~~ ligand is an avidin or a strepavidin molecule.
44. (Currently Amended) The method of claim 40, wherein the reagent-reacted ~~molecule~~ protein is further isolated by removing substantially all of the non-bound ~~molecule~~ proteins from the ligand-bound ~~molecule~~ proteins.
45. (Currently Amended) The method of claim 40, wherein the non-bound ~~molecule~~ protein is removed by washing.
46. (Currently Amended) The method of claim 40, wherein the reagent-reacted ~~molecule~~ is further isolated by cleaving step ~~the cleavable chemical moiety of the cell membrane impermeable reagent under a condition that does not denature the lumen-exposed molecule and does not dissociate the ligand from the binding domain after removing a non-bound molecule~~ protein.
- 47-48. (Cancelled)
49. (Currently Amended) The method of claim 46, wherein the reagent-reacted ligand-bound lumen-exposed ~~molecule~~ protein is further isolated by elution from the binding domain and the ligand.
50. (Cancelled)

51. (Currently Amended) A method of isolating an organ-specific or tissue-specific ~~molecule~~ protein that is exposed on a luminal surface of an arteriole, a capillary or a vein comprising the following steps:
- (a) providing a cell membrane impermeable reagent comprising three domains
    - (i) a first domain comprising an active moiety capable of covalently and non-specifically binding to a molecule expressed on the luminal surface of a cell lining a perfusable space *in situ* or *in vivo*,
    - (ii) a second domain comprising a biotin binding domain, and
    - (iii) a third domain comprising a disulfide moiety situated between the first and second domains linking the first domain to the second domain; and
  - (b) administering the cell membrane impermeable reagent into a lumen of an artery, a arteriole, a capillary or a vein in an intact organ or an intact animal to react the cell membrane impermeable reagent with a ~~molecule~~ protein expressed on the luminal surface; and
  - (d) isolating the reagent-reacted ~~molecule~~ protein by contacting the reagent-reacted ~~molecule~~ protein with an immobilized avidin or streptavidin molecule;
  - (e) removing substantially all of the non-immobilized ~~molecules~~ proteins; and
  - (f) cleaving the cleavable chemical moiety of the reagent that reacted with the lumen exposed protein under a condition that does not denature the lumen-exposed protein.

52-55. (Cancelled)

56. (Currently Amended) The method of claim 44 10, wherein the enzyme is selected from the group consisting of horseradish peroxidase, alkaline phosphatase,  $\beta$ -galactosidase, and acetylcholinesterase.

57. (Currently Amended) The method of claim 44 10, wherein the bioluminescent moiety is selected from the group consisting of luciferase, luciferin, and aequorin.

58. (Previously Presented) The method of claim 10, wherein the radionucleotide is selected from the group consisting of H-3, S-35, I-125, I-131, P-32, Y-90, Re-188, At-211, and Bi-212.

59. (Previously Presented) The method of claim 10, wherein the paramagnetic moiety is selected from the group consisting of Cr, V, Mn, Fe, Co, Ni, Cu, La, Ce, Pr, Nd, Pm, Sm, Eu, Gd, Tb, Dy, Ho, Er, Tm, Yb, and Lu.

60. (Previously Presented) The method of claim 10, wherein the fluorescent moiety is selected from the group consisting of umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride, and phycoerythrin.

61. (Previously Presented) The method of claim 51, further comprising the step of cleaving the cleavable chemical moiety of the cell membrane impermeable reagent under a condition that does not dissociate said immobilized avidin or streptavidin molecule from said biotin binding domain.

62. (New) The method of claim 51, further comprising the step of comparing the reagent-reacted proteins from different organs or tissues to identify the organ-specific or tissue-specific protein.

63. (New) The method of claim 52 wherein the organ-specific or tissue-specific protein is detected in only one tissue.